



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Mr. Gray S. Nunnelly  
Chief Executive Officer  
Pure Company  
2045 Preisker Lane  
Santa Maria, California 93454

JUL - 3 1997

Re: K971278  
Trade Name: Pure Company Fluid Delivery System  
Regulatory Class: I  
Product Code: EIA  
Dated: April 4, 1997  
Received: April 7, 1997

Dear Mr. Nunnelly:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to 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). 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 (Pre-market Approval), 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 895. A substantially equivalent determination assumes compliance with the current Good Manufacturing Practice requirement, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic (QS) inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531

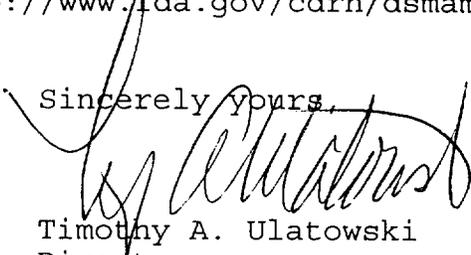
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through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 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 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4618. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

Sincerely yours,



Timothy A. Ulatowski  
Director  
Division of Dental, Infection Control  
and General Hospital Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

510(k) Number: K971278

Device Name: Pure Fluid Delivery System

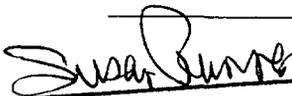
Indications For Use: ---

The Pure Fluid Delivery System is a dual reservoir system that isolates a dental operator from municipal or rural water supplies. It can also be used to introduce bottled solutions that are commonly used in dental procedures through the dental unit. The Pure Fluid Delivery System also has a .01 Micron air filter that permits air purging of the dental unit with higher quality air.

The operation of the the Fluid Delivery System is facilitated by a rotary switch on the control face that permits switching between the contents of the two bottles being used. There is a built-in pressure relief valve that limits pressure in the bottles to 40psi as a safety feature.

The control face panel also has a quick disconnect that allows the solutions in the bottle selected to be used in an external dental device.

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

 Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)  
Division of Dental, Infection Control,  
and General Hospital Devices

510(k) Number K971278

Prescription Use   
(Per 21 CFR 801.109)

OR

Over the Counter Use