



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JUN 4 1999

Mr. Miguel A. Arrizon
PARI Respiratory Equipment, Inc.
13800 Hull Street Road
Midlothian, VA 23112

Re: K983308
RC Cornet, Model #44F50
Regulatory Class: II (two)
Product Code: 73 BWF
Dated: March 4, 1999
Received: March 8, 1999

Dear Mr. Arrizon:

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 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). 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.

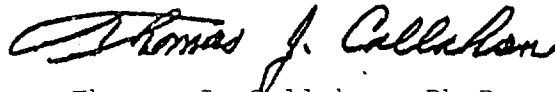
If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, 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 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

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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) 594-4648. 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/dsma/dsmamain.html>".

Sincerely yours,

A handwritten signature in black ink, reading "Thomas J. Callahan". The signature is fluid and cursive, with the first name "Thomas" being the most prominent part.

Thomas J. Callahan, Ph.D.
Director
Division of Cardiovascular,
Respiratory, and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

May 28, 1999

Emil Wang
FDA, CDRH
Document Mail Center (HFZ-401)
9200 Corporate Boulevard
Rockville, MD 20850 USA

RE: K983308, Revised Indications For Use Statement

Dear Mr. Wang,

Following is the Revised Indications For Use Statement.

PARI
Respiratory Equipment, Inc.

IN USA
13800 Hull Street Road
Midlothian, VA 23112 USA
1-800-FAST-NEB (327-863)
Tel: 804.639.7235
Fax: 804.639.7242

IN CANADA
3120 Glen Erin Drive
Unit 10
Mississauga, Ontario L5L 1
1-800-825-1637
Tel: 905.828.9119
Fax: 905.828.5436



The PARI RC Cornet Mucus Clearing Device is a Positive Expiratory Pressure PEP Device and it was designed to exercise patient's lungs and to improve secretion clearance. The device was designed to work with PARI LC Nebulizers.

If you have any questions please contact me at extension 237.

Best Regards,

Miguel A. Arrizon
Regulatory Affairs Manager

(Division Sign-Off)
Division of Cardiovascular, Respiratory,
and Neurological Devices
510(k) Number K983308

prescriptions use

OTC

Specialists in effective
drug inhalation
Spécialistes de
l'inhalothérapie efficace
Especializados en
inhaloterapia eficaz