K992553

APR 12 2000

EXHIBIT 7

510(k) Summary

Submitted by: Daniel J. Manelli
Farkas & Manelli, P.L.L.C.
2000 M Street, NW (Suite 700)
Washington, DC 20036

On behalf of Dormer Laboratories, Inc.
510(k) Submission: IQ Chamber
July 28, 1999

The product is a plastic tray with shallow indentations or chambers for use as a holding device to place allergens or allergen mixes in contact with the surface of the skin during allergen patch tests. .

This product is for use only by health care professionals; it is not intended for OTC use. It contains materials that pose no health hazard when used according to directions. The device is substantially equivalent to various marketed allergen and vaccine delivery systems including Hill Top Chambers (K811365)

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

APR 12 2000

Dormer Laboratories, Inc.
C/O Mr. Daniel J. Manelli
Farkas & Manelli, PLLC
2000 M Street, N.W., 7th Floor
Washington, D.C. 20036-3307

Re: K992553
Trade Name: IQ Chamber
Regulatory Class: I
Product Code: LDH
Dated: February 22, 2000
Received: February 24, 2000

Dear Mr. Manelli:

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 (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 Good Manufacturing Practice for Medical Devices: General (GMP) regulation (21 CFR Part 820) and that, through periodic GMP 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.

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) 594-4690. 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

EXHIBIT 5

510(k) Number (if known): K992553

Device Name: IQ Chamber

Indications For Use:

The IQ Chamber is an allergen delivery system which places allergens or allergen mixes in contact with the surface of the skin for the purpose of performing patch testing.

(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 K992553

Prescription Use ✓ OR Over-The-Counter Use _____
(Per 21 CFR 801.109)