



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Windmere Corporation  
c/o Mr. Richard C. Lanzillotto  
Vice President  
North American Technical Services Corporation  
30 Northport Road  
Sound Beach, New York 11789-1734

JUN 13 1997

\*Re: K970590  
Health Zone/Healthy Vibes Mini Rechargeable  
Infrared Massager, Model KM-213H  
K970592  
Health Zone/Healthy Vibes Infrared  
Massager Set, Model KM-201  
Regulatory Class: II  
Product Codes: ILY and ISA  
Dated: April 29, 1997  
Received: May 5, 1997

Dear Mr. Lanzillotto:

We have reviewed your Section 510(k) notification of intent to market the devices referenced above and we have determined these devices are 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 devices, 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.

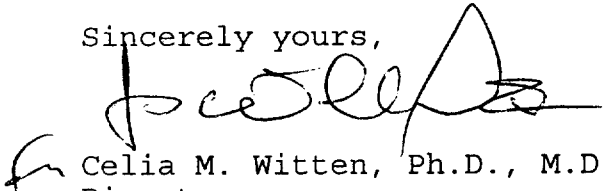
If your devices are classified (see above) into either class II (Special Controls) or class III (Premarket Approval), they may be subject to such additional controls. Existing major regulations affecting your devices can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially 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 devices 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 devices as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your devices to legally marketed predicate devices results in a classification for your devices and thus, permits your devices to proceed to the market.

If you desire specific advice for your devices 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) 594-4659. Additionally, for questions on the promotion and advertising of your devices, 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,

  
Celia M. Witten, Ph.D., M.D.  
Director  
Division of General and  
Restorative Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosures

K970592/A<sup>2</sup>

# NATS Corp.

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Engineering, Regulatory and Consulting Services  
Product and Process Management

REF : 510K K970592 WINDMERE MODEL KM-201

IN KEEPING WITH OUR CONVERSATION, ALL CLAIMS WILL BE DELETED EXCEPT AS FOLLOWS:

- RELIEVES MINOR MUSCLE ACHES AND PAINS
- DRY HEAT THERAPY FOR BODY SURFACES

Over-the-Counter Use X

[Signature]  
(Division Sign-Off)

Division of General Restorative Devices

510(k) Number

K970592