



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

MAR 8 2002

ENSD Corporation
c/o Mr. Kevin Walls, RAC
Principal Consultant
Regulatory Insight, Inc.
13 Red Fox Lane
Littleton, CO 80127

Re: K020094

Trade/Device Name: EN 600-658 Standard Wheelchair
EN 600-666 Standard Wheelchair
EN 600-664 Standard Wheelchair
EN 600-690 Extra Wide Wheelchair
EN 600-651 Aluminum Light Weight Wheelchair-29 lbs
EN 600-663 Light Weight Wheelchair-32 lbs
EN 600-665 Standard Wheelchair
EN 600-670 Reclinable Wheelchair

Regulation Number: 890.3850

Regulation Name: Wheelchair, mechanical

Regulatory Class: I

Product Code: IOR

Dated: January 10, 2002

Received: January 11, 2002

Dear Mr. Walls:

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.

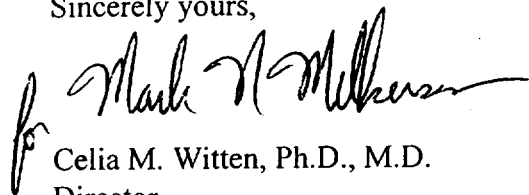
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish 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. Page 2 -

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 and additionally 21 CFR Part 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 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 Part 807.97). Other general information on your responsibilities under the Act may be obtained 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/dsma/dsmamain.html>

Sincerely yours,

A handwritten signature in black ink, appearing to read "Celia M. Witten", with a stylized initial "f" to the left.

Celia M. Witten, Ph.D., M.D.
Director

Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number (if known): K020094

Device Name: EN 600-658 Standard Wheelchair
 EN 600-666 Standard Wheelchair
 EN 600-664 Standard Wheelchair
 EN 600-690 Extra Wide Wheelchair
 EN 600-651 Alum. Light Weight Wheelchair-29lbs
 EN 600-663 Light Weight Wheelchair-32lbs
 EN 600-665 Standard Wheelchair
 EN 600-670 Reclinable Wheelchair

Indications for Use: The EN 600-658, EN 600-666, EN 600-664, EN 600-690, EN 600-651, EN 600-663, EN 600-665 and EN 600-670 Mechanical Wheelchairs are manually operated devices with wheels that are intended for medical purposes to provide mobility to persons restricted to a sitting position.

PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON ANOTHER PAGE IF NEEDED

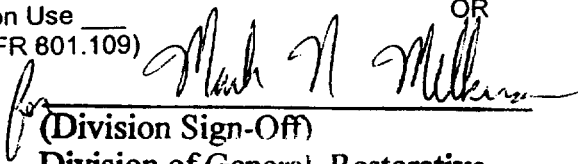
Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of General, Restorative and
Neurological Devices
510(k) Number _____

Prescription Use _____
(Per 21 CFR 801.109)

OR

Over the Counter Use X


(Division Sign-Off)
Division of General, Restorative
and Neurological Devices

510(k) Number K020094