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### Device Listing Database

**Proprietary Device Name:** BIO-MAT  
**Common/Generic Device Name:** PAD, HEATING, POWERED  
**Classification Name:** PAD, HEATING, POWERED  
**Product Code:** [IRT](#)  
**Device Class:** 2  
**Regulation Number:** [890.5740](#)  
**Medical Specialty:** Physical Medicine  
**Owner/Operator:** [CHARNTECH ELECTRONIC](#)  
**Owner/Operator Number:** 9056316  
**Registered Establishment Name:** [RICHWAY INTL... INC.](#)  
**Establishment Registration Number:** 2954299  
**Date of Listing:** 05/24/99  
**Listing Status:** Active  
**Establishment Operations:** Initial Distributor

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Center for Devices and Radiological Health / CDRH



FEB - 8 2008

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

RichWay International, Inc  
% Mr. LeRoy Klima  
Consultant  
1314 South King Street, Suite 520  
Honolulu, Hawaii 96814

Re: K072534  
Trade/Device Name: Bio-Mat Mattress  
Regulation Number: 21 CFR 890.5740  
Regulation Name: Powered heating pad, Infrared lamp  
Regulatory Class: Class II  
Product Code: IRT  
Dated: January 03, 2008  
Received: January 08, 2008

Dear Mr. Klima:

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](#).

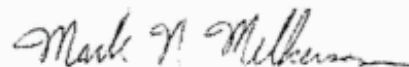
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's 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 – Mr. LeRoy Klima

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), please contact the Center for Devices and Radiological Health's (CDRH's) Office of Compliance at (240) 276-0120. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding postmarket surveillance, please contact CDRH's Office of Surveillance and Biometric's (OSB's) Division of Postmarket Surveillance at (240) 276-3474. For questions regarding the reporting of device adverse events (Medical Device Reporting (MDR)), please contact the Division of Surveillance Systems at (240) 276-3464. You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,



Mark N. Melkerson  
Director  
Division of General, Restorative  
and Neurological Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

## Indications for Use

510(k) Number \_\_\_\_\_

**Device Name:** Bio-Mat Mattress

### Indications for Use:

The Bio-Mat is indicated for the temporary relief of minor muscle and joint pain and stiffness; the temporary relief of joint pain associated with arthritis; the temporary relief of muscle spasms, minor sprains and strains, and minor muscular back pain; the relaxation of muscles; and the temporary increase of local circulation where applied.

Prescription Use \_\_\_\_\_  
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use X \_\_\_\_\_  
(21 CFR 801 Subpart C)



\_\_\_\_\_  
(Division Sign-Off)

**Division of General, Restorative,  
and Neurological Devices**

510(k) Number 16072534