

**MAC Medical**  
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Millstadt, IL 62260  
Phone (618) 476-3550  
Fax (618) 476-3337  
Date prepared: May 25, 2008  
Contact: Dennis Cooper

**1. Identification of the Device:**

Proprietary-Trade Name: SWC/DWC (Various Models) Warming Cabinets  
Classification Name: Warmer, ~~irrigation solution cabinet~~  
Product Code ~~HC~~ LDO  
Common/Usual Name: ~~Irrigation Warmer~~

**2. Equivalent legally marketed devices:** K993797, Entermics Medical Systems Fluid Warming Cabinet

**3. Indications for Use (intended use)** These warming cabinets are designed to store, warm, and maintain the recommended temperature of irrigation solutions prior to use.

**4. Description of the Devices:** These devices are 120 or 220 V Ac powered warming cabinets designed to store, warm, and maintain the recommended temperature of irrigation solutions prior to use. The particular cabinet configuration can be deduced from its model number:

SWC = Single Warming Cabinet.

DWC = Dual Warming Cabinet.

G = Glass Door

The temperature ranges and capacities vary according to model but all have:

Adjustable Temperature Range

Digital Control

Dual Display

Push Button Operation

Rapid Warm Time

High Accuracy.

**5. Safety and Effectiveness, comparison to predicate device.** The results of bench, user, and standards testing indicates that the new device is as safe and effective as the predicate devices.

## 6. Substantial Equivalence Chart

Characteristic	K993797, Enthermics Medical Systems Fluid Warming Cabinet	SWC/DWC (Various Models) Warming Cabinets
Intended Use:	Warming Irrigation Solutions	SAME
User Interface	Digital Display	SAME
Power Source	AC Line	SAME
Door opening	Left or right hand	SAME
Door style	Glass	Glass or Stainless
Construction	Stainless Steel	SAME
Safety	UL Listed	SAME

## 7. Conclusion

After analyzing bench and standards testing data, it is the conclusion of MAC Medical that the SWC/DWC (Various Models) Warming Cabinets are as safe and effective as the predicate devices, have few technological differences, and have no new indications for use, thus rendering them substantially equivalent to the predicate devices.



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Mac Medical  
C/O Mr. Daniel Kamm  
Regulatory Engineer  
Kamm & Associates  
P.O. Box 7007  
Deerfield, Illinois 60015

**JUN 13 2008**

Re: K080609  
Trade/Device Name: SWC/DWC Warming Cabinets  
Regulation Number: Unclassified  
Generic Name: Device, General Medical, Warming Cabinet  
Regulatory Class: Unclassified  
Product Code: LDQ  
Dated: May 30, 2008  
Received: June 2, 2008

Dear Mr. Kamm:

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.

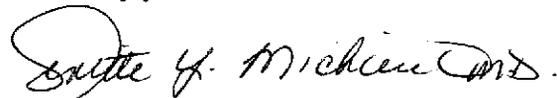
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish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

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 Office of Compliance at (240) 276-0115. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). 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 (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,



Chiu Lin, Ph.D.  
Director  
Division of Anesthesiology, General Hospital,  
Infection Control and Dental Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

## Indications for Use

510(k) Number (if known): K 080609

Device Name: SWC/DWC (Various Models) Warming Cabinets

### Indications For Use:

These warming cabinets are designed to store, warm, and maintain the recommended temperature of irrigation solutions prior to use.

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

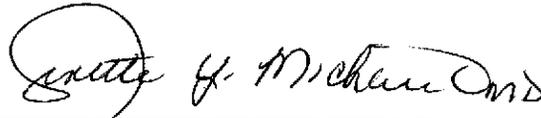
AND/OR

Over-The-Counter Use X  
(21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

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Concurrence of CDRH, Office of Device Evaluation (ODE)



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(Division Sign-Off)

Division of Anesthesiology, General Hospital  
Infection Control, Dental Devices

510(k) Number: K 080609