

K974454

P1091

#### 4. Safety and Effectiveness 510(k) Summary

DEC 18 1997

Date Prepared: November 20, 1997

Submitted By:

Tucson Medical Corporation  
3941 East 29th Street, Suite 601  
Tucson, AZ 85711  
Phone: (520) 512 - 1100, Fax: (520) 512-8019  
Contact Person: Douglas Bueschel

Trade name: ClearField Anti-Fog Sterile Wipe, Product Number 300-004.

Common name: External Lens Anti-Fog Solution.

Classification name: Endoscope Accessory (per 21 CFR section 876.1500).

Predicate Device: ClearField Anti-Fog Sterile Wipe, 510(k) Number K904871/B.

Device Description: Sterile cloth pre-saturated with 5cc of Anti-Fog solution. Single Use.

Intended Use: Intended for use as an agent to keep external viewing surfaces clear of water vapor and condensation.

Substantial equivalence is based upon equivalent technological characteristics of the solution, cloth and ink of this device compared to the predicate device.



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

DEC 18 1997

Mr. Jim Patko  
Operations Manager  
Tucson Medical Corporation  
3941 East 29<sup>th</sup> Street, Suite 601  
Tucson, Arizona 85711

Re: K974454  
Clear-Field Anti-Fog Sterile Wipe (Product Number 300-004)  
Dated: November 20, 1997  
Received: November 25, 1997  
Regulatory class: II  
21 CFR §876.1500/Product code: 78 KOG

Dear Mr. Patko:

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 Current Good Manufacturing Practice requirement, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS 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) 594-4613. 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,

Lillian Yin, Ph.D.  
Director, Division of Reproductive,  
Abdominal, Ear, Nose and Throat,  
and Radiological Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

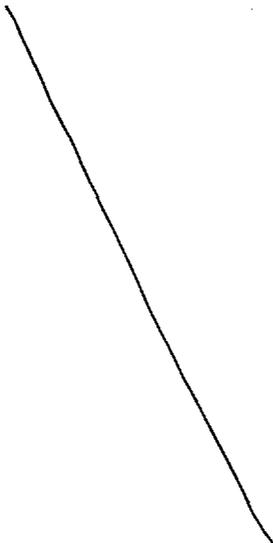
Enclosure

510(k) Number (if known): K974454

Device Name: ClearField Anti-Fog Sterile Wipes

Indications For Use:

ClearField Anti-Fog Sterile Wipe is intended for use as an agent to keep external viewing surfaces clear of water vapor and condensation.



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

Concurrence of CDRH, Office of Device Evaluation (ODE)

*Robert R. Smith*  
(Division Sign-Off)  
Division of Reproductive, Abdominal, ENT,  
and Radiological Devices  
510(k) Number K974454

Prescription Use   
(Per 21 CFR 801.109)

OR

Over-The-Counter Use

(Optional Format 1-2-96)