



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

NOV 6 1998

Frank E. Platko, Ph.D.  
Vice President, Research & Development  
Propper Manufacturing Company, Incorporated  
36-04 Skillman Avenue  
Long Island City, New York 11101

Re: K973585  
Trade Name: Propper EO LINE™ Ethylene Oxide  
Sterilization Integrator  
Regulatory Class: II  
Product Code: JOJ  
Dated: September 3, 1998  
Received: September 4, 1998

Dear Dr. Platko:

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 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). 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.

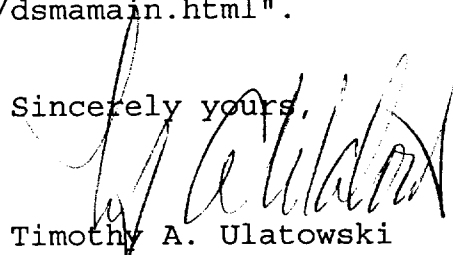
If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, 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) 594-4692. 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" (21CFR 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/dsma/dsmamain.html>".

Sincerely yours,



Timothy A. Ulatowski  
Director  
Division of Dental, Infection Control,  
and General Hospital Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

K973585

# Propper EO LINE™ Ethylene Oxide Sterilization Integrator

### Indications For Use:

Propper EO LINE™ is indicated to be an ethylene oxide sterilization integrator for use in EO sterilizers with 10% ethylene oxide : 90% HCFC gas mixtures.

Its function is to monitor the exposure conditions for EO sterilization and to ensure that the four critical parameters: EO gas concentration, temperature, humidity and exposure time have been met.

The performance of Propper EO LINE™ Integrator was comparable to SteriGage® EO Integrator when tested in a Joslyn BIER vessel.

All testing was conducted with 10% EO : 90% HCFC gas mixtures.

**(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 Dental, Infection Control,  
and General Health Services

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

Prescription Use \_\_\_\_\_  
(Per 21 CFR 801.109)

OR

### Over-The-Counter Use