



JUL 17 1998

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Mr. Paul Schiller
Worldwide Product Manager
Eastman Kodak Company
343 State Street
Rochester, New York 14650

Re: K973430
Trade Name: Kodak Pro-Medical Digital Camera System
Regulatory Class: II
Product Code: GCJ
Dated: March 12, 1998
Received: March 16, 1998

Dear Mr. Schiller:

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.

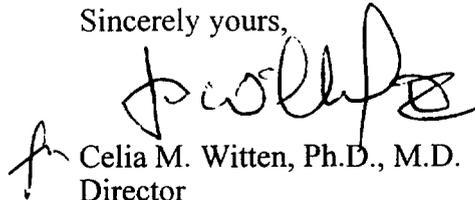
If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially 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.

Page 2 - Mr. Paul Schiller

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) 594-4595. 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,



Celia M. Witten, Ph.D., M.D.
Director
Division of General and
Restorative Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number (if known): 973430

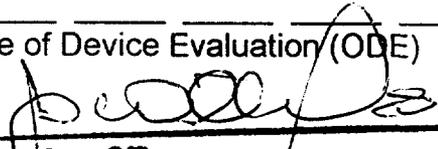
Device Name: KODAK Pro-Medical Digital Camera System

Indications For Use:

The camera is intended to digitally capture images for micro-surgery, pathology, micro-biology, ophthalmology and general surgery applications. Upon capturing the image, it is stored in the memory of the PCMCIA Card in the camera. User may utilize the audio function of the camera to record patient, exam or any other pertinent identification information. The Camera is connected to the computer via the high speed serial bus (IEEE 1394) cable interface. The images are viewed, selected and then downloaded using Twain Compliant Driver for PCs and Adobe Photoshop Plug-in Software for MACINTOSH based systems. The user will be able to annotate and format the images and add information prior to transmission and/or archiving. The camera is capable of both lossy and lossless data compression. The operator is instructed in the manual to utilize only lossless data compression for medical applications.

(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 Devices
510(k) Number 973430

Prescription Use (Per 21 CFR 801.109)

OR

Over-The-Counter Use

(Optional Format 1-2-96)