FDA 510(k) Premarket Notification

K033821 KODAK DRYVIEW 8900 LASER IMAGER MAMMOGRAPHY ACCESSORY

CLEARED

Feb 20, 2004
73 days to decision

K033821 · Product code: **LMC** · Radiology
Source: https://www.510kdatabase.net/k033821/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Multi Format, Radiological (LMC)
Date received	Dec 9, 2003
Decision date	Feb 20, 2004
Days to decision	73 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Eastman Kodak Company**
Location	Mchenry, IL, US
Contact	STEPHEN SLAVENS
Website	http://www.kodak.com
510(k) history	238 submissions · 238 cleared · 1977-2006

Eastman Kodak Company is a diversified imaging and materials manufacturer headquartered in McHenry, US. The company has a long history in advanced materials, chemicals, and imaging technologies. Eastman Kodak maintains a significant regulatory history in medical imaging devices. The company received FDA 510(k) clearances from total submissions, with clearances spanning from 1977 to 2006. The company's cleared devices focused primarily on radiology and medical imaging systems, including digital radiography systems, picture archiving and communication systems (PACS), and re...