

K926404 KODAK X-OMATIC IDENTIFICATION CAMERA, MODEL 4 & 4LMar 4, 1993
73 days to decisionK926404 · Product code: **JAC** · Radiology
Source: <https://www.510kdatabase.net/k926404/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Film Marking, Radiographic (JAC)
Date received	Dec 21, 1992
Decision date	Mar 4, 1993
Days to decision	73 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Eastman Kodak Company
Location	Mchenry, IL, US
Contact	NORMAN H GEIL
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...

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k926404/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 18, 2026