

**K060550 KODAK DIRECTVIEW DR 3000 SYSTEM 64KW/80KW
GENERATORS, MODELS 6551345, 6551386**Mar 31, 2006
29 days to decisionK060550 · Product code: **KPR** · Radiology
Source: <https://www.510kdatabase.net/k060550/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, X-ray, Stationary (KPR)
Date received	Mar 2, 2006
Decision date	Mar 31, 2006
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Eastman Kodak Company
Location	Mchenry, IL, US
Contact	CHRISTINE E EHMANN
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...

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Device record: <https://www.510kdatabase.net/k060550/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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