

**K974107 KODAK DIGITAL SCIENCE QUALITY CONTROL
WORKSTATION (QCW)**Jan 21, 1998
82 days to decisionK974107 · Product code: **LMD** · Radiology
Source: <https://www.510kdatabase.net/k974107/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Digital Image Communications, Radiological (LMD)
Date received	Oct 31, 1997
Decision date	Jan 21, 1998
Days to decision	82 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Eastman Kodak Company
Location	Mchenry, IL, US
Contact	NANCY BUTCHER
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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