FDA 510(k) Premarket Notification

K770837 SCREEN, FINE, LANES, KODAK

CLEARED

May 26, 1977
20 days to decision

K770837 · Product code: **EAM** · Radiology
Source: https://www.510kdatabase.net/k770837/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screen, Intensifying, Radiographic (EAM)
Date received	May 6, 1977
Decision date	May 26, 1977
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Eastman Kodak Company**
Location	Mchenry, IL, US
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...