FDA 510(k) Premarket Notification

K840134 CAMTEX 1000

CLEARED

Mar 16, 1984
72 days to decision

K840134 · Product code: **JAA** · Radiology
Source: https://www.510kdatabase.net/k840134/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	Jan 4, 1984
Decision date	Mar 16, 1984
Days to decision	72 days
Third-party review	No

APPLICANT

Company	**Digital Imaging Co. of America**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1984-1984