FDA 510(k) Premarket Notification

K864837 MODIFIED MAINFRAME OF MODEL 902 MOBILE C-ARM

CLEARED

Jan 30, 1987
52 days to decision

K864837 · Product code: **JAA** · Radiology
Source: https://www.510kdatabase.net/k864837/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	Dec 9, 1986
Decision date	Jan 30, 1987
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Oec-Diasonics, Inc.**
Location	Salt Lake City, UT, US
Contact	JOHN W TOLHURST
510(k) history	8 submissions · 8 cleared · 1985-1993