

K864837 MODIFIED MAINFRAME OF MODEL 902 MOBILE C-ARMJan 30, 1987
52 days to decisionK864837 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k864837/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026