FDA 510(k) Premarket Notification

K874934 UNIVERSAL EXAMINATION TABLE

CLEARED

Jan 19, 1988
46 days to decision

K874934 · Product code: **KXJ** · Radiology
Source: https://www.510kdatabase.net/k874934/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Radiologic (KXJ)
Date received	Dec 4, 1987
Decision date	Jan 19, 1988
Days to decision	46 days
Third-party review	No

APPLICANT

Company	**Imatron, Inc.**
Location	South San Fransisco, CA, US
Contact	J. A CODUTO
510(k) history	11 submissions · 11 cleared · 1983-2000