FDA 510(k) Premarket Notification

K871827 OMEGA III ANGIOGRAPHIC TABLE, ROTATION & STEPPER

CLEARED

Jun 19, 1987
39 days to decision

K871827 · Product code: **KXJ** · Radiology
Source: https://www.510kdatabase.net/k871827/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Radiologic (KXJ)
Date received	May 11, 1987
Decision date	Jun 19, 1987
Days to decision	39 days
Third-party review	No

APPLICANT

Company	**General Electric Co.**
Location	Mchenry, IL, US
Contact	LARRY KROGER
510(k) history	254 submissions · 254 cleared · 1976-2011

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k871827/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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