FDA 510(k) Premarket Notification

K862120 SCX SYSTEM

K862120 · Product code: **JAA** · Radiology
Source: https://www.510kdatabase.net/k862120/

CLEARED

Jun 16, 1986
13 days to decision

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	Jun 3, 1986
Decision date	Jun 16, 1986
Days to decision	13 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k862120/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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