FDA 510(k) Premarket Notification

K770094 FLUORICON 500 IMAGE INTENSIFIERS

CLEARED

Jan 21, 1977
4 days to decision

K770094 · Product code: **EAM** · Radiology
Source: https://www.510kdatabase.net/k770094/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screen, Intensifying, Radiographic (EAM)
Date received	Jan 17, 1977
Decision date	Jan 21, 1977
Days to decision	4 days
Third-party review	No

APPLICANT

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