

K770094 FLUORICON 500 IMAGE INTENSIFIERSJan 21, 1977
4 days to decisionK770094 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k770094/>; 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 17, 2026