

K862414 FUJI CR IMAGING PLATEFeb 26, 1987
246 days to decisionK862414 · Product code: **EAM** · Radiology
Source: <https://www.510kdatabase.net/k862414/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screen, Intensifying, Radiographic (EAM)
Date received	Jun 25, 1986
Decision date	Feb 26, 1987
Days to decision	246 days
Third-party review	No

APPLICANT

Company	Parexel Intl. Corp.
Location	Mchenry, IL, US
Contact	SAYIGH, PHD
510(k) history	14 submissions · 14 cleared · 1984-1991

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k862414/>; 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 21, 2026