

**K893510 CRONEX QUANTA V RARE EARTH INTENSIFYING
SCREEN**Jun 23, 1989
49 days to decisionK893510 · Product code: **EAM** · Radiology
Source: <https://www.510kdatabase.net/k893510/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screen, Intensifying, Radiographic (EAM)
Date received	May 5, 1989
Decision date	Jun 23, 1989
Days to decision	49 days
Third-party review	No

APPLICANT

Company	E.I. Dupont DE Nemours & Co., Inc.
Location	Mchenry, IL, US
Contact	KENNETH L WOODLIN
510(k) history	253 submissions · 252 cleared · 1976-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893510/> 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 14, 2026