

**K845030 CRONEX SR342-QUANTA FAST DETAIL-INTENSI-
SCREENS**Feb 26, 1985
61 days to decisionK845030 · Product code: **EAM** · Radiology
Source: <https://www.510kdatabase.net/k845030/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screen, Intensifying, Radiographic (EAM)
Date received	Dec 27, 1984
Decision date	Feb 26, 1985
Days to decision	61 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k845030/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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