FDA 510(k) Premarket Notification

K845030 CRONEX SR342-QUANTA FAST DETAIL-INTENSI-SCREENS

CLEARED

Feb 26, 1985
61 days to decision

K845030 · 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