FDA 510(k) Premarket Notification

K893510 CRONEX QUANTA V RARE EARTH INTENSIFYING SCREEN

CLEARED

Jun 23, 1989
49 days to decision

K893510 · 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