FDA 510(k) Premarket Notification

K770574 CRONEX QUANTA III INTENSIFYING SCREENS

CLEARED

Apr 5, 1977
8 days to decision

K770574 · Product code: **EAM** · Radiology
Source: https://www.510kdatabase.net/k770574/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screen, Intensifying, Radiographic (EAM)
Date received	Mar 28, 1977
Decision date	Apr 5, 1977
Days to decision	8 days
Third-party review	No

APPLICANT

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