FDA 510(k) Premarket Notification

K933161 ULTRA-VISION SUPER RAPID INTENSIFYING SCREEN

CLEARED

Oct 7, 1993
100 days to decision

K933161 · Product code: **EAM** · Radiology
Source: https://www.510kdatabase.net/k933161/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screen, Intensifying, Radiographic (EAM)
Date received	Jun 29, 1993
Decision date	Oct 7, 1993
Days to decision	100 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dupont Medical Products**
Location	Wilmington, DE, US
Contact	JEAN BARTLETT
510(k) history	28 submissions · 28 cleared · 1991-1995