FDA 510(k) Premarket Notification

CLEARED

K900935 DU PONT LINX LASER IMAGER W/THE LINX NETWORKING

May 15, 1990
76 days to decision

K900935 · Product code: **LMD** · Radiology
Source: https://www.510kdatabase.net/k900935/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Digital Image Communications, Radiological (LMD)
Date received	Feb 28, 1990
Decision date	May 15, 1990
Days to decision	76 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