

**K900935 DU PONT LINX LASER IMAGER W/THE LINX NETWORKING**May 15, 1990  
76 days to decisionK900935 · Product code: **LMD** · Radiology  
Source: <https://www.510kdatabase.net/k900935/>**SUBMISSION DETAILS**

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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**

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Company	<b>E.I. Dupont DE Nemours &amp; Co., Inc.</b>
Location	Mchenry, IL, US
Contact	KENNETH L WOODLIN
510(k) history	253 submissions · 252 cleared · 1976-1996

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900935/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 14, 2026