

**K020243 LASERPRO 16**Mar 5, 2002  
41 days to decisionK020243 · Product code: **LMA** · Radiology  
Source: <https://www.510kdatabase.net/k020243/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Digitizer, Image, Radiological (LMA)
Date received	Jan 23, 2002
Decision date	Mar 5, 2002
Days to decision	41 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Eradlink, Inc.</b>
Location	Penn Valley, CA, US
Contact	ALBERT J KOUBA
510(k) history	2 submissions · 2 cleared · 2002-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020243/> 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 18, 2026