FDA 510(k) Premarket Notification

K020243 LASERPRO 16

CLEARED

Mar 5, 2002
41 days to decision

K020243 · Product code: **LMA** · Radiology
Source: https://www.510kdatabase.net/k020243/

SUBMISSION DETAILS

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

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