FDA 510(k) Premarket Notification

K110840 LIAC

CLEARED

Apr 17, 2012
389 days to decision

K110840 · Product code: **IYE** · Radiology
Source: https://www.510kdatabase.net/k110840/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Mar 25, 2011
Decision date	Apr 17, 2012
Days to decision	389 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Sordina S.P.A.**
Location	Ormond Beach, FL, US
Contact	BERTHOIN CLAUDE
510(k) history	1 submissions · 1 cleared · 2012-2012