

K110840 LIACApr 17, 2012
389 days to decisionK110840 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k110840/> Data sourced from FDA 510(k) public records (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. - Generated May 17, 2026