

**K934303 RFA A-300**Aug 9, 1994  
342 days to decisionK934303 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k934303/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Sep 1, 1993
Decision date	Aug 9, 1994
Days to decision	342 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Scanditronix, Inc.</b>
Location	Newburyport, MA, US
Contact	LENNART WARD
510(k) history	7 submissions · 7 cleared · 1984-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k934303/> 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 20, 2026