

**K030162 LAWRENCE CSERION, MODEL CS-1**Mar 28, 2003  
71 days to decisionK030162 · Product code: **KXK** · Radiology  
Source: <https://www.510kdatabase.net/k030162/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Source, Brachytherapy, Radionuclide (KXK)
Date received	Jan 16, 2003
Decision date	Mar 28, 2003
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Isoray, Inc.</b>
Location	Richland, WA, US
Contact	DAVID SWANBERG
510(k) history	3 submissions · 3 cleared · 2003-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030162/> 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 17, 2026