FDA 510(k) Premarket Notification

K030162 LAWRENCE CSERION, MODEL CS-1

CLEARED

Mar 28, 2003
71 days to decision

K030162 · Product code: **KXK** · Radiology
Source: https://www.510kdatabase.net/k030162/

SUBMISSION DETAILS

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

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