FDA 510(k) Premarket Notification

K091602 MUCHECK V8

CLEARED

Jun 18, 2009
15 days to decision

K091602 · Product code: **IYE** · Radiology
Source: https://www.510kdatabase.net/k091602/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jun 3, 2009
Decision date	Jun 18, 2009
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Oncology Data Systems, Inc.**
Location	Oklahoma City, OK, US
Contact	VINCE RUMINER
510(k) history	4 submissions · 4 cleared · 2001-2010