FDA 510(k) Premarket Notification

K092822 CAPRI APPLICATOR

CLEARED

Nov 3, 2009
50 days to decision

K092822 · Product code: **JAQ** · Radiology
Source: https://www.510kdatabase.net/k092822/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Applicator, Radionuclide, Remote-controlled (JAQ)
Date received	Sep 14, 2009
Decision date	Nov 3, 2009
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vivaray, Inc.**
Location	Portola Valley, CA, US
Contact	GEORGE HERMANN
510(k) history	1 submissions · 1 cleared · 2009-2009