

**K092822 CAPRI APPLICATOR**Nov 3, 2009  
50 days to decisionK092822 · 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	<b>Vivaray, Inc.</b>
Location	Portola Valley, CA, US
Contact	GEORGE HERMANN
510(k) history	1 submissions · 1 cleared · 2009-2009

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092822/> 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