FDA 510(k) Premarket Notification

K093092 ODYSSEY WORKSTATION

CLEARED

Jan 21, 2010
112 days to decision

K093092 · Product code: **DQK** · Cardiovascular
Source: https://www.510kdatabase.net/k093092/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Oct 1, 2009
Decision date	Jan 21, 2010
Days to decision	112 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stereotaxis, Inc.**
Location	St. Louis, MO, US
Contact	DENNIS POZZO
510(k) history	28 submissions · 28 cleared · 2002-2026