FDA 510(k) Premarket Notification

K122888 CDR SYSTEMS PRECISION PATIENT POSITIONING SYSTEM

CLEARED

Apr 30, 2013
222 days to decision

K122888 · Product code: **IYE** · Radiology
Source: https://www.510kdatabase.net/k122888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Sep 20, 2012
Decision date	Apr 30, 2013
Days to decision	222 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cdr Systems, Inc.**
Location	Calgary, CA
Contact	CARL DENIS
510(k) history	1 submissions · 1 cleared · 2013-2013