

**K122888 CDR SYSTEMS PRECISION PATIENT POSITIONING SYSTEM**

Apr 30, 2013  
222 days to decision

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

**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k122888/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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