FDA 510(k) Premarket Notification

K111862 VIEWRAY SYSTEM

CLEARED

May 22, 2012
327 days to decision

K111862 · Product code: **IYE** · Radiology
Source: https://www.510kdatabase.net/k111862/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jun 30, 2011
Decision date	May 22, 2012
Days to decision	327 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Viewray, Incorporated**
Location	Oakwood Village, OH, US
Contact	JANICE BROWNLEE
510(k) history	6 submissions · 6 cleared · 2011-2021