FDA 510(k) Premarket Notification

K123371 ALIGNRT PLUS

CLEARED

Jan 8, 2013
68 days to decision

K123371 · Product code: **IYE** · Radiology
Source: https://www.510kdatabase.net/k123371/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Nov 1, 2012
Decision date	Jan 8, 2013
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vision RT Limited**
Location	London, GB
Contact	NORMAN SMITH
510(k) history	3 submissions · 3 cleared · 2006-2013