

K123371 ALIGNRT PLUSJan 8, 2013
68 days to decisionK123371 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k123371/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 17, 2026