

**K131313 ACTIVE BREATHING COORDINATOR**Aug 7, 2013  
92 days to decisionK131313 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k131313/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	May 7, 2013
Decision date	Aug 7, 2013
Days to decision	92 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aktina Medical Corp.</b>
Location	Congers, NY, US
Contact	TONY SPACCAROTELLA
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/k131313/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 17, 2026