

**K131907 RAYPEX 6**Jan 31, 2014  
219 days to decisionK131907 · Product code: **LQY** · Dental  
Source: <https://www.510kdatabase.net/k131907/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Locator, Root Apex (LQY)
Date received	Jun 26, 2013
Decision date	Jan 31, 2014
Days to decision	219 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dentsply International, Inc.</b>
Location	York, PA, US
Contact	HELEN LEWIS
510(k) history	36 submissions · 36 cleared · 2006-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131907/> 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 June 2, 2026