

**K083057 I-MAX TOUCH**Jun 2, 2009  
231 days to decisionK083057 · Product code: **MUH** · Radiology  
Source: <https://www.510kdatabase.net/k083057/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Extraoral Source, Digital (MUH)
Date received	Oct 14, 2008
Decision date	Jun 2, 2009
Days to decision	231 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Owandy</b>
Location	Ormond Beach, FL, US
Contact	CLAUDE D BERTHOIN
510(k) history	6 submissions · 6 cleared · 2006-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083057/> 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 16, 2026