

**K053010 VISTADENT AT COMPLETE, MODEL 3.1**

Nov 22, 2005  
27 days to decision

K053010 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k053010/>

**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Oct 26, 2005
Decision date	Nov 22, 2005
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Dentsply Intl., Inc.</b>
Location	York, PA, US
Contact	HELEN LEWIS
510(k) history	12 submissions · 12 cleared · 2005-2014

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k053010/> 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 18, 2026