

**K990983 MODIFICATION TO IMTEC SENDAX MDI**Apr 13, 1999  
20 days to decisionK990983 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k990983/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Mar 24, 1999
Decision date	Apr 13, 1999
Days to decision	20 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Intec Corp.</b>
Location	Ardmore, OK, US
Contact	M.K. PATTERSON
510(k) history	28 submissions · 28 cleared · 1991-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990983/> 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