

**K000889 SENOJ IMPLANT SYSTEM**Apr 27, 2001  
403 days to decisionK000889 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k000889/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Mar 20, 2000
Decision date	Apr 27, 2001
Days to decision	403 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Senoj Biocare, Inc.</b>
Location	San Diego, CA, US
Contact	PENNY WOLLUM
510(k) history	1 submissions · 1 cleared · 2001-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000889/> 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 July 2, 2026