

**K981052 CRESCO TI IMPLANT CONCEPT**Dec 18, 2001  
1366 days to decisionK981052 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k981052/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Mar 23, 1998
Decision date	Dec 18, 2001
Days to decision	1366 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Cresco TI Systems AB</b>
Location	Kristianstad, SE
Contact	PER E NILSSON
510(k) history	2 submissions · 2 cleared · 2001-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k981052/> 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 1, 2026