

**K022859 ABUTMENT AND PROSTHETIC ACCESSORIES TO THE ITI DENTAL IMPLANT SYSTEM**

Sep 19, 2002  
22 days to decision

K022859 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k022859/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Aug 28, 2002
Decision date	Sep 19, 2002
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Straumann USA</b>
Location	Waltham, MA, US
Contact	LINDA JALBERT
510(k) history	41 submissions · 41 cleared · 1996-2016

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

Device record: <https://www.510kdatabase.net/k022859/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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