

**K071439 ZIMMER PATIENT-SPECIFIC ABUTMENT, INTERNAL
HEX, TITANIUM**

Sep 20, 2007
120 days to decision

K071439 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k071439/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	May 23, 2007
Decision date	Sep 20, 2007
Days to decision	120 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Zimmer Dental, Inc.
Location	Carlsbad, CA, US
Contact	ERIN MCVEY
510(k) history	40 submissions · 40 cleared · 2005-2017

510k Database - www.510kdatabase.net
 Device record: <https://www.510kdatabase.net/k071439/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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