

**K072363 NANOTITE PREVAIL,CERTAIN,PARELLEL
WALLED,TAPERED,EXTERNAL HEX**Jan 31, 2008
161 days to decisionK072363 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k072363/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Aug 23, 2007
Decision date	Jan 31, 2008
Days to decision	161 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biomet 3i, Inc.
Location	Palm Beach Gardens, FL, US
Contact	DIANA TAYLOR
510(k) history	13 submissions · 13 cleared · 2007-2011

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072363/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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