

K073368 5.0 X 5.0MM DENTAL IMPLANT AND 6.0 X 5.0 DENTAL IMPLANTOct 10, 2008
315 days to decisionK073368 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k073368/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Nov 30, 2007
Decision date	Oct 10, 2008
Days to decision	315 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Bicon, LLC
Location	Boston, MA, US
Contact	VINCENT J MORGAN
510(k) history	5 submissions · 5 cleared · 2008-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073368/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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