

**K012473 VIDENT HEXTOP, SYNOCTA IN-CERAM  
BLANK/COPING MANUFACTURED BY VIDENT**Sep 11, 2002  
405 days to decisionK012473 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k012473/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Aug 2, 2001
Decision date	Sep 11, 2002
Days to decision	405 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Vident</b>
Location	Baldwin Park, CA, US
Contact	BARBARA J LEWANDOWSKI
510(k) history	36 submissions · 36 cleared · 1990-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012473/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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