

**K013672 BRUGES DISTRACTION - ANCHORING -  
OSTEOSYNTHESIS SYSTEM**Apr 8, 2002  
153 days to decision

K013672 · Product code: JEY · Dental

Source: <https://www.510kdatabase.net/k013672/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	Nov 6, 2001
Decision date	Apr 8, 2002
Days to decision	153 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Azary Technologies, LLC</b>
Location	Huntington, CT, US
Contact	JOSEPH M AZARY
510(k) history	1 submissions · 1 cleared · 2002-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013672/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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