

**K923419 TISMESH SYSTEM**Apr 8, 1993  
269 days to decisionK923419 · Product code: **JEY** · DentalSource: <https://www.510kdatabase.net/k923419/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	Jul 13, 1992
Decision date	Apr 8, 1993
Days to decision	269 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ti-Mesh, Inc.</b>
Location	Las Vegas, NV, US
Contact	MARY P MORGAN
510(k) history	3 submissions · 3 cleared · 1992-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923419/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 3, 2026