

**K023360 LEFORTE SYSTEM BONE PLATE**Oct 25, 2002  
18 days to decisionK023360 · Product code: **KTW** · Orthopedic  
Source: <https://www.510kdatabase.net/k023360/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Single Component (KTW)
Date received	Oct 7, 2002
Decision date	Oct 25, 2002
Days to decision	18 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Jeil Medical Corporation</b>
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	53 submissions · 53 cleared · 2002-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023360/> 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 May 20, 2026