

**K023851 PRECIMED HIP SCREW SYSTEM**Jan 28, 2003  
70 days to decisionK023851 · Product code: **KTT** · Orthopedic  
Source: <https://www.510kdatabase.net/k023851/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Nov 19, 2002
Decision date	Jan 28, 2003
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Precimed, Inc.</b>
Location	Chester Springs, PA, US
Contact	BARBARA LYONS
510(k) history	2 submissions · 2 cleared · 2003-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023851/> 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