

**K032634 NORMED COMPRESSION BONE SCREW SYSTEM**Oct 6, 2003  
41 days to decisionK032634 · Product code: **JDS** · Orthopedic  
Source: <https://www.510kdatabase.net/k032634/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nail, Fixation, Bone (JDS)
Date received	Aug 26, 2003
Decision date	Oct 6, 2003
Days to decision	41 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Noviace Corporation</b>
Location	Paramus, NJ, US
Contact	ALBERT ENAYATI
510(k) history	2 submissions · 2 cleared · 2003-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032634/> 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