

**K063248 ARCUATE VERTEBRAL AUGMENTATION SYSTEM**Mar 13, 2007  
138 days to decisionK063248 · Product code: **NDN** · Orthopedic  
Source: <https://www.510kdatabase.net/k063248/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cement, Bone, Vertebroplasty (NDN)
Date received	Oct 26, 2006
Decision date	Mar 13, 2007
Days to decision	138 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Sofamor Danek</b>
Location	Memphis, TN, US
Contact	CHRISTINE SCIFERT
510(k) history	154 submissions · 147 cleared · 2002-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063248/> 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 16, 2026