

**K024060 MODIFICATION TO ASNIS III CANNULATED SCREW SYSTEM**Dec 20, 2002  
11 days to decisionK024060 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k024060/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Screw, Fixation, Bone (HWC)
Date received	Dec 9, 2002
Decision date	Dec 20, 2002
Days to decision	11 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Howmedica Osteonics Corp.</b>
Location	Allendale, NJ, US
Contact	KAREN ARIEMMA
510(k) history	288 submissions · 288 cleared · 1999-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k024060/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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