

K862157 OMC ORTHOMET CANNULATED SCREWAug 29, 1986
85 days to decision

K862157 · Orthopedic

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jun 5, 1986
Decision date	Aug 29, 1986
Days to decision	85 days
Third-party review	No

APPLICANT

Company	Orthomet, Inc.
Location	Plymouth, MN, US
Contact	PRO TEM
510(k) history	60 submissions · 41 cleared · 1986-1995

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Device record: <https://www.510kdatabase.net/k862157/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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