

**K883941 PROFORMA HIP SYSTEM**Nov 30, 1988  
72 days to decisionK883941 · Product code: **JDI** · Orthopedic  
Source: <https://www.510kdatabase.net/k883941/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Sep 19, 1988
Decision date	Nov 30, 1988
Days to decision	72 days
Third-party review	No

**APPLICANT**

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Company	<b>Orthomet, Inc.</b>
Location	Plymouth, MN, US
Contact	DENNIS H CRANE
510(k) history	60 submissions · 41 cleared · 1986-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883941/>, 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 21, 2026