

**K930275 PRECISION OSTEOLOCK ACETABULAR COMPONENT**

Sep 1, 1994  
590 days to decision

K930275 · Product code: **JDI** · Orthopedic  
Source: <https://www.510kdatabase.net/k930275/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Jan 19, 1993
Decision date	Sep 1, 1994
Days to decision	590 days
Third-party review	No

**APPLICANT**

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Company	<b>Howmedica Corp.</b>
Location	Mchenry, IL, US
Contact	ROBERT E SMITH
510(k) history	373 submissions · 325 cleared · 1976-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k930275/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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