

**K902395 MODIFIED AML + ACETABULAR CUP SYSTEM**Aug 24, 1990  
86 days to decisionK902395 · Product code: **JDI** · Orthopedic  
Source: <https://www.510kdatabase.net/k902395/>**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	May 30, 1990
Decision date	Aug 24, 1990
Days to decision	86 days
Third-party review	No

**APPLICANT**

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Company	<b>Depuy, Inc.</b>
Location	Mchenry, IL, US
Contact	KENNETH G ROBERTS
510(k) history	303 submissions · 239 cleared · 1976-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902395/>; 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 20, 2026