

K800894 DUPUY CEMENT RESTRICTOR TRIALS & INSTR.May 2, 1980
14 days to decisionK800894 · Product code: **JDI** · Orthopedic
Source: <https://www.510kdatabase.net/k800894/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Apr 18, 1980
Decision date	May 2, 1980
Days to decision	14 days
Third-party review	No

APPLICANT

Company	Depuy, Inc.
Location	Mchenry, IL, US
510(k) history	303 submissions · 239 cleared · 1976-2005

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Device record: <https://www.510kdatabase.net/k800894/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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