

K830750 CEMENTED 101 32 FEMORAL COMPONENTMay 2, 1983
112 days to decisionK830750 · Product code: **JDI** · Orthopedic
Source: <https://www.510kdatabase.net/k830750/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Jan 10, 1983
Decision date	May 2, 1983
Days to decision	112 days
Third-party review	No

APPLICANT

Company	Intermedics Orthopedics
Location	Mchenry, IL, US
510(k) history	108 submissions · 82 cleared · 1982-1997

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Device record: <https://www.510kdatabase.net/k830750/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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