

K812672 SELF-CENTERING HIPOct 8, 1981
16 days to decisionK812672 · Product code: **KWY** · Orthopedic
Source: <https://www.510kdatabase.net/k812672/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Hemi-, Femoral, Metal/polymer, Cemented Or Uncemented (KWY)
Date received	Sep 22, 1981
Decision date	Oct 8, 1981
Days to decision	16 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/k812672/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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