

K790996 C.C.F. FEMORAL DEBRIDEMENT PROBEJun 27, 1979
29 days to decisionK790996 · Product code: **HTO** · Orthopedic
Source: <https://www.510kdatabase.net/k790996/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reamer (HTO)
Date received	May 29, 1979
Decision date	Jun 27, 1979
Days to decision	29 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/k790996/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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