

K800363 PARKE-DAVIS TOTAL KNEE PROSTHESISMar 10, 1980
19 days to decisionK800363 · Product code: **KTZ** · Orthopedic
Source: <https://www.510kdatabase.net/k800363/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Caliper (KTZ)
Date received	Feb 20, 1980
Decision date	Mar 10, 1980
Days to decision	19 days
Third-party review	No

APPLICANT

Company	Warner-Lambert Co.
Location	Mchenry, IL, US
510(k) history	50 submissions · 50 cleared · 1979-2003

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Device record: <https://www.510kdatabase.net/k800363/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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