FDA 510(k) Premarket Notification

K800363 PARKE-DAVIS TOTAL KNEE PROSTHESIS

CLEARED

Mar 10, 1980
19 days to decision

K800363 · 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