FDA 510(k) Premarket Notification

K062960 CONSERVE FEMORAL RESURFACING COMPONENT

CLEARED

Dec 1, 2006
63 days to decision

K062960 · Product code: **KXA** · Orthopedic
Source: https://www.510kdatabase.net/k062960/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Hip, Femoral, Resurfacing (KXA)
Date received	Sep 29, 2006
Decision date	Dec 1, 2006
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Wrightmedicaltechnologyinc**
Location	Arlington, TN, US
Contact	THERESA LEISTER
510(k) history	302 submissions · 291 cleared · 1993-2023