FDA 510(k) Premarket Notification

K082673 CONSERVE PRESSFIT FEMORAL COMPONENT

CLEARED

Dec 12, 2008
88 days to decision

K082673 · Product code: **KXA** · Orthopedic
Source: https://www.510kdatabase.net/k082673/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Femoral, Resurfacing (KXA)
Date received	Sep 15, 2008
Decision date	Dec 12, 2008
Days to decision	88 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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k082673/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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