FDA 510(k) Premarket Notification

K050699 MODIFICATION TO: VENTED CEMENT RESTRICTOR, MODEL 60102-000-001/004

CLEARED

Apr 15, 2005
28 days to decision

K050699 · Product code: **JDK** · Orthopedic
Source: https://www.510kdatabase.net/k050699/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Hip, Cement Restrictor (JDK)
Date received	Mar 18, 2005
Decision date	Apr 15, 2005
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Amedica Corp.**
Location	Salt Lake City, UT, US
Contact	ROBERT M WOLFARTH
510(k) history	16 submissions · 16 cleared · 2002-2018

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k050699/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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