

K022615 CENTERPULSE, SPINE-TECH DIVISION CEMENT RESTRICTOR

Oct 29, 2002
83 days to decision

K022615 · Product code: **JDK** · Orthopedic
Source: <https://www.510kdatabase.net/k022615/>

SUBMISSION DETAILS

Decision	Substantially Equivalent - U
Submission type	Special
Device classification	Prosthesis, Hip, Cement Restrictor (JDK)
Date received	Aug 7, 2002
Decision date	Oct 29, 2002
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Centerpulse Spine-Tech, Inc.
Location	Minneapolis, MN, US
Contact	ANGELA BYLAND
510(k) history	4 submissions · 2 cleared · 2002-2004

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k022615/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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