FDA 510(k) Premarket Notification

K022374 CADENCE SPINAL FIXATION SYSTEM

CLEARED

Sep 24, 2002
64 days to decision

K022374 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k022374/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Jul 22, 2002
Decision date	Sep 24, 2002
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sulzer Spine-Tech**
Location	Minneapolis, MN, US
Contact	KRISTYN M BENSON
510(k) history	7 submissions · 7 cleared · 1999-2002