FDA 510(k) Premarket Notification

K063083 MODIFICATION TO CD HORIZON SPINAL SYSTEM

CLEARED

Nov 6, 2006
27 days to decision

K063083 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k063083/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Oct 10, 2006
Decision date	Nov 6, 2006
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Sofamor Danek, Inc.**
Location	Memphis, TN, US
Contact	EDWARD S CHIN
510(k) history	99 submissions · 89 cleared · 2000-2025