FDA 510(k) Premarket Notification

K031780 VERTE-STACK SPINAL SYSTEM

CLEARED

Jul 30, 2003
50 days to decision

K031780 · Product code: **MQP** · Orthopedic
Source: https://www.510kdatabase.net/k031780/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Jun 10, 2003
Decision date	Jul 30, 2003
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Sofamor Danek**
Location	Memphis, TN, US
Contact	RICHARD TREHARNE
510(k) history	154 submissions · 147 cleared · 2002-2021