FDA 510(k) Premarket Notification

K001844 MODIFICATION TO CENTAUR SPINAL SYSTEM

CLEARED

Jul 19, 2000
30 days to decision

K001844 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k001844/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jun 19, 2000
Decision date	Jul 19, 2000
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Howmedica Osteonics Corp.**
Location	Allendale, NJ, US
Contact	MARY-CATHERINE DILLON
510(k) history	288 submissions · 288 cleared · 1999-2020