FDA 510(k) Premarket Notification

K032059 MODIFICATION TO MACS MODULAR ANTERIOR CONSTRUCT SYSTEM

CLEARED

Jul 25, 2003
23 days to decision

K032059 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k032059/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jul 2, 2003
Decision date	Jul 25, 2003
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Aesculap, Inc.**
Location	Burlingame, CA, US
Contact	GEORG KELLER
510(k) history	207 submissions · 201 cleared · 1991-2025