FDA 510(k) Premarket Notification

K032371 TITANIUM MESH SYSTEM

CLEARED

Feb 12, 2004
195 days to decision

K032371 · Product code: **MQP** · Orthopedic
Source: https://www.510kdatabase.net/k032371/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Aug 1, 2003
Decision date	Feb 12, 2004
Days to decision	195 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Encore Medical, L.P.**
Location	Austin, TX, US
Contact	KIMBERLY L PRUITT
510(k) history	81 submissions · 81 cleared · 2002-2026