FDA 510(k) Premarket Notification

K041407 TITANIUM MESH IMPLANT

CLEARED

Dec 29, 2004
216 days to decision

K041407 · Product code: **MQP** · Orthopedic
Source: https://www.510kdatabase.net/k041407/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	May 27, 2004
Decision date	Dec 29, 2004
Days to decision	216 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Interpore Cross Intl.**
Location	Irvine, CA, US
Contact	WENDY SPIELBERGER
510(k) history	39 submissions · 38 cleared · 1998-2005