FDA 510(k) Premarket Notification

K051781 GRAFTCAGE TLX

CLEARED

Dec 16, 2005
168 days to decision

K051781 · Product code: **MQP** · Orthopedic
Source: https://www.510kdatabase.net/k051781/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Jul 1, 2005
Decision date	Dec 16, 2005
Days to decision	168 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Osteotech, Inc.**
Location	San Mateo, CA, US
Contact	CHRISTOPHER TALBOT
510(k) history	24 submissions · 21 cleared · 1985-2008