FDA 510(k) Premarket Notification

K120724 PIONEER LATERAL PLATE SYSTEM

CLEARED

May 7, 2012
59 days to decision

K120724 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k120724/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Mar 9, 2012
Decision date	May 7, 2012
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Pioneer Surgical Technology, Inc.**
Location	Marquette, MI, US
Contact	SARAH MCINTYRE
510(k) history	26 submissions · 26 cleared · 2010-2025