FDA 510(k) Premarket Notification

K051665 SYNTHES VECTRA-T SYSTEM

CLEARED

Sep 9, 2005
79 days to decision

K051665 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k051665/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jun 22, 2005
Decision date	Sep 9, 2005
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Synthes Spine**
Location	Paoli, PA, US
Contact	SUSAN LEWANDOWSKI
510(k) history	36 submissions · 32 cleared · 1995-2012