FDA 510(k) Premarket Notification

K000742 SYNTHES ANTERIOR CSLP SYSTEM

CLEARED

Mar 29, 2000
22 days to decision

K000742 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k000742/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Mar 7, 2000
Decision date	Mar 29, 2000
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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