FDA 510(k) Premarket Notification

K010515 MODIFICATION TO SYNERGY SPINAL SYSTEM

CLEARED

Mar 22, 2001
28 days to decision

K010515 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k010515/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Feb 22, 2001
Decision date	Mar 22, 2001
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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