FDA 510(k) Premarket Notification

K990845 MODIFICATION TO PARTS NUMBERED CGT-SL AND CGT-SL-15

CLEARED

Apr 13, 1999
29 days to decision

K990845 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k990845/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Mar 15, 1999
Decision date	Apr 13, 1999
Days to decision	29 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Advanced Spine Fixation Systems, Inc.**
Location	Washington, DC, US
Contact	GREG HOLLAND
510(k) history	16 submissions · 7 cleared · 1989-2000