FDA 510(k) Premarket Notification

K983706 MODIFICATION OF TOWNLEY PEDICLE SCREW PLATING SYSTEM

CLEARED

Nov 12, 1998
22 days to decision

K983706 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k983706/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Oct 21, 1998
Decision date	Nov 12, 1998
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sofamor Danek USA,Inc.**
Location	Memphis, TN, US
Contact	RICHARD W TREHARNE
510(k) history	41 submissions · 26 cleared · 1995-2000