FDA 510(k) Premarket Notification

K895439 ACROMED BEADED SUBLAMINAR WIRE

CLEARED

Nov 22, 1989
78 days to decision

K895439 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k895439/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Sep 5, 1989
Decision date	Nov 22, 1989
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Acromed Corp.**
Location	Cleveland, OH, US
Contact	ZANG, RN
510(k) history	41 submissions · 22 cleared · 1984-1998