FDA 510(k) Premarket Notification

K923703 KANEDA ANTERIOR SPINAL MULTISEGMENTAL FIXATION DEV

CLEARED

May 20, 1993
300 days to decision

K923703 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k923703/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jul 24, 1992
Decision date	May 20, 1993
Days to decision	300 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k923703/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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