FDA 510(k) Premarket Notification

K820660 A HOOK DRIVER

CLEARED

Mar 19, 1982
9 days to decision

K820660 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k820660/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Mar 10, 1982
Decision date	Mar 19, 1982
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**Twin City Surgical, Inc.**
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1981-1987