

K820660 A HOOK DRIVERMar 19, 1982
9 days to decisionK820660 · Product code: **KWP** · Orthopedic
Source: <https://www.510kdatabase.net/k820660/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (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

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Device record: <https://www.510kdatabase.net/k820660/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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