

K800729 PROBEApr 8, 1980
7 days to decisionK800729 · Product code: **HXB** · Orthopedic
Source: <https://www.510kdatabase.net/k800729/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe (HXB)
Date received	Apr 1, 1980
Decision date	Apr 8, 1980
Days to decision	7 days
Third-party review	No

APPLICANT

Company	Stainless Mfg., Inc.
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1980-1984

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Device record: <https://www.510kdatabase.net/k800729/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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