

K800730 RETROGRADE KNIFEApr 8, 1980
7 days to decisionK800730 · Product code: **HXP** · Orthopedic
Source: <https://www.510kdatabase.net/k800730/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Bending Or Contouring (HXP)
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/k800730/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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