FDA 510(k) Premarket Notification

K800729 PROBE

CLEARED

Apr 8, 1980
7 days to decision

K800729 · 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