

K802951 I.M PROBESDec 17, 1980
27 days to decisionK802951 · Product code: **HXB** · Orthopedic
Source: <https://www.510kdatabase.net/k802951/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe (HXB)
Date received	Nov 20, 1980
Decision date	Dec 17, 1980
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Hogan & Hartson
Location	Mchenry, IL, US
510(k) history	26 submissions · 25 cleared · 1980-1998

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Device record: <https://www.510kdatabase.net/k802951/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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