FDA 510(k) Premarket Notification

K802951 I.M PROBES

CLEARED

Dec 17, 1980
27 days to decision

K802951 · 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