FDA 510(k) Premarket Notification

K770184 BI-CENTRIC ENDOPROSTHESES

CLEARED

Feb 8, 1977
12 days to decision

K770184 · Product code: **JDG** · Orthopedic
Source: https://www.510kdatabase.net/k770184/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Femoral Component, Cemented, Metal (JDG)
Date received	Jan 27, 1977
Decision date	Feb 8, 1977
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Howmedica Corp.**
Location	Mchenry, IL, US
510(k) history	373 submissions · 325 cleared · 1976-1998