FDA 510(k) Premarket Notification

K830750 CEMENTED 101 32 FEMORAL COMPONENT

CLEARED

May 2, 1983
112 days to decision

K830750 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k830750/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Jan 10, 1983
Decision date	May 2, 1983
Days to decision	112 days
Third-party review	No

APPLICANT

Company	**Intermedics Orthopedics**
Location	Mchenry, IL, US
510(k) history	108 submissions · 82 cleared · 1982-1997