FDA 510(k) Premarket Notification

K883941 PROFORMA HIP SYSTEM

CLEARED

Nov 30, 1988
72 days to decision

K883941 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k883941/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Sep 19, 1988
Decision date	Nov 30, 1988
Days to decision	72 days
Third-party review	No

APPLICANT

Company	**Orthomet, Inc.**
Location	Plymouth, MN, US
Contact	DENNIS H CRANE
510(k) history	60 submissions · 41 cleared · 1986-1995