FDA 510(k) Premarket Notification

K930275 PRECISION OSTEOLOCK ACETABULAR COMPONENT

CLEARED

Sep 1, 1994
590 days to decision

K930275 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k930275/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Jan 19, 1993
Decision date	Sep 1, 1994
Days to decision	590 days
Third-party review	No

APPLICANT

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