FDA 510(k) Premarket Notification

K800894 DUPUY CEMENT RESTRICTOR TRIALS & INSTR.

CLEARED

May 2, 1980
14 days to decision

K800894 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k800894/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Apr 18, 1980
Decision date	May 2, 1980
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Depuy, Inc.**
Location	Mchenry, IL, US
510(k) history	303 submissions · 239 cleared · 1976-2005