FDA 510(k) Premarket Notification

K843909 TITANIUM HIP 1019

CLEARED

Nov 15, 1984
43 days to decision

K843909 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k843909/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Oct 3, 1984
Decision date	Nov 15, 1984
Days to decision	43 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k843909/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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