FDA 510(k) Premarket Notification

K861979 AML II TITANIUM ACETABULUM PROSTHESIS

CLEARED

Jan 15, 1987
239 days to decision

K861979 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k861979/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	May 21, 1986
Decision date	Jan 15, 1987
Days to decision	239 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k861979/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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