FDA 510(k) Premarket Notification

K902395 MODIFIED AML + ACETABULAR CUP SYSTEM

CLEARED

Aug 24, 1990
86 days to decision

K902395 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k902395/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	May 30, 1990
Decision date	Aug 24, 1990
Days to decision	86 days
Third-party review	No

APPLICANT

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