FDA 510(k) Premarket Notification

K903636 OSTEONICS DRM ACETABULAR COMPONENT

CLEARED

Oct 30, 1990
78 days to decision

K903636 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k903636/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Aug 13, 1990
Decision date	Oct 30, 1990
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Osteonics Corp.**
Location	Mchenry, IL, US
Contact	PAT KRAMER
510(k) history	178 submissions · 136 cleared · 1980-2000