FDA 510(k) Premarket Notification

K911052 NEXUS(TM) FEMORAL HIP STEM

CLEARED

Jun 17, 1991
101 days to decision

K911052 · Product code: **JDI** · Orthopedic
Source: https://www.510kdatabase.net/k911052/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Mar 8, 1991
Decision date	Jun 17, 1991
Days to decision	101 days
Third-party review	No

APPLICANT

Company	**Dow Corning Wright**
Location	Mchenry, IL, US
Contact	MICKI ROY
510(k) history	74 submissions · 52 cleared · 1979-1994