FDA 510(k) Premarket Notification

K862136 MODIFIED WHITESIDE POSTERIOR STABILIZED TOTAL KNEE

CLEARED

Jul 14, 1986
40 days to decision

K862136 · Product code: **HSX** · Orthopedic
Source: https://www.510kdatabase.net/k862136/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Non-constrained, Cemented, Metal/polymer (HSX)
Date received	Jun 4, 1986
Decision date	Jul 14, 1986
Days to decision	40 days
Third-party review	No

APPLICANT

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