FDA 510(k) Premarket Notification

K902538 WHITESIDE ORTHOLOC MODULAR TIBIAL AUGMENTATION

CLEARED

Jul 6, 1990
30 days to decision

K902538 · Product code: **HRY** · Orthopedic
Source: https://www.510kdatabase.net/k902538/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Jun 6, 1990
Decision date	Jul 6, 1990
Days to decision	30 days
Third-party review	No

APPLICANT

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