FDA 510(k) Premarket Notification

K881779 WHITESIDE ORTHOLOC II UNICONDYLAR KNEE SYSTEM

CLEARED

Aug 31, 1988
127 days to decision

K881779 · Product code: **HSX** · Orthopedic
Source: https://www.510kdatabase.net/k881779/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Non-constrained, Cemented, Metal/polymer (HSX)
Date received	Apr 26, 1988
Decision date	Aug 31, 1988
Days to decision	127 days
Third-party review	No

APPLICANT

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