FDA 510(k) Premarket Notification

K810540 KNEE PROSTHESIS

CLEARED

Apr 7, 1981
39 days to decision

K810540 · Product code: **HRY** · Orthopedic
Source: https://www.510kdatabase.net/k810540/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Feb 27, 1981
Decision date	Apr 7, 1981
Days to decision	39 days
Third-party review	No

APPLICANT

Company	**Cintor Orthopaedic Div.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1980-1981