FDA 510(k) Premarket Notification

K790350 POSTERIOR CRUCIATE CONDYLAR TIBIAL

CLEARED

Mar 6, 1979
13 days to decision

K790350 · Product code: **HRY** · Orthopedic
Source: https://www.510kdatabase.net/k790350/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Feb 21, 1979
Decision date	Mar 6, 1979
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Howmedica Corp.**
Location	Mchenry, IL, US
510(k) history	373 submissions · 325 cleared · 1976-1998