FDA 510(k) Premarket Notification

K860133 MODIFIED SYNATOMIC VARIABLE FIT TIBIAL PLATEAU

CLEARED

Mar 10, 1986
55 days to decision

K860133 · Product code: **HRY** · Orthopedic
Source: https://www.510kdatabase.net/k860133/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Jan 14, 1986
Decision date	Mar 10, 1986
Days to decision	55 days
Third-party review	No

APPLICANT

Company	**Depuy, Inc.**
Location	Mchenry, IL, US
Contact	STEVE WENTWORTH
510(k) history	303 submissions · 239 cleared · 1976-2005

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k860133/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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