FDA 510(k) Premarket Notification

K860594 SYNATOMIC REVISION FEMORAL PROSTHESIS

CLEARED

Mar 14, 1986
23 days to decision

K860594 · Product code: **JWH** · Orthopedic
Source: https://www.510kdatabase.net/k860594/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Feb 19, 1986
Decision date	Mar 14, 1986
Days to decision	23 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/k860594/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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