FDA 510(k) Premarket Notification

K862125 ARIZONA CONDYLAR TIBIAL PLATEAU

CLEARED

Jun 27, 1986
24 days to decision

K862125 · Product code: **HRY** · Orthopedic
Source: https://www.510kdatabase.net/k862125/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Jun 3, 1986
Decision date	Jun 27, 1986
Days to decision	24 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k862125/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026