FDA 510(k) Premarket Notification

K812672 SELF-CENTERING HIP

CLEARED

Oct 8, 1981
16 days to decision

K812672 · Product code: **KWY** · Orthopedic
Source: https://www.510kdatabase.net/k812672/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Hemi-, Femoral, Metal/polymer, Cemented Or Uncemented (KWY)
Date received	Sep 22, 1981
Decision date	Oct 8, 1981
Days to decision	16 days
Third-party review	No

APPLICANT

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