FDA 510(k) Premarket Notification

K896669 WEBER PERMALOCK

CLEARED

Feb 6, 1990
71 days to decision

K896669 · Product code: **LZO** · Orthopedic
Source: https://www.510kdatabase.net/k896669/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/ceramic/polymer, Cemented Or Non-porous, Uncemented (LZO)
Date received	Nov 27, 1989
Decision date	Feb 6, 1990
Days to decision	71 days
Third-party review	No

APPLICANT

Company	**Allo Pro Corp.**
Location	Mchenry, IL, US
Contact	K PIERPOINT
510(k) history	16 submissions · 15 cleared · 1981-1990