FDA 510(k) Premarket Notification

K900051 MOORE TYPE ENDO PROSTHESIS

CLEARED

Oct 3, 1990
273 days to decision

K900051 · Product code: **KWY** · Orthopedic
Source: https://www.510kdatabase.net/k900051/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Hemi-, Femoral, Metal/polymer, Cemented Or Uncemented (KWY)
Date received	Jan 3, 1990
Decision date	Oct 3, 1990
Days to decision	273 days
Third-party review	No

APPLICANT

Company	**Lima Intl. Corp.**
Location	E Providence, RI, US
Contact	POLLONIO-LISTER
510(k) history	8 submissions · 7 cleared · 1989-1994