

**K900051 MOORE TYPE ENDO PROSTHESIS**Oct 3, 1990  
273 days to decisionK900051 · 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	<b>Lima Intl. Corp.</b>
Location	E Providence, RI, US
Contact	POLLONIO-LISTER
510(k) history	8 submissions · 7 cleared · 1989-1994

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900051/>, 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 22, 2026