

**K896669 WEBER PERMALOCK**Feb 6, 1990  
71 days to decisionK896669 · 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	<b>Allo Pro Corp.</b>
Location	Mchenry, IL, US
Contact	K PIERPOINT
510(k) history	16 submissions · 15 cleared · 1981-1990

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896669/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 22, 2026