

K923909 CEMENT RESTRICTOR FOR PROFORMA AND PERFECTA TOTAL

Nov 4, 1992
92 days to decision

K923909 · Product code: **LZN** · Orthopedic
Source: <https://www.510kdatabase.net/k923909/>

SUBMISSION DETAILS

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Cement Obturator (LZN) |
| Date received | Aug 4, 1992 |
| Decision date | Nov 4, 1992 |
| Days to decision | 92 days |
| Third-party review | No |
| Summary / Statement | Statement |

APPLICANT

| | |
|----------------|---|
| Company | Orthomet, Inc. |
| Location | Plymouth, MN, US |
| Contact | DAVID A CANNISTRACI |
| 510(k) history | 60 submissions · 41 cleared · 1986-1995 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k923909/> 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 21, 2026