

**K081860 ZIMMER SEGMENTAL SYSTEM VARIABLE STIFFNESS
STEM EXTENSIONS AND INTERCALARY SEGMENTS**Sep 26, 2008
87 days to decisionK081860 · Product code: **LZO** · Orthopedic
Source: <https://www.510kdatabase.net/k081860/>**SUBMISSION DETAILS**

| | |
|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Prosthesis, Hip, Semi-constrained, Metal/ceramic/polymer, Cemented Or Non-porous, Uncemented (LZO) |
| Date received | Jul 1, 2008 |
| Decision date | Sep 26, 2008 |
| Days to decision | 87 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Zimmer, Inc. |
| Location | Warsaw, IN, US |
| Contact | TONI KINGSLEY |
| Website | https://www.zimmerbiomet.com |
| 510(k) history | 373 submissions · 352 cleared · 1976-2026 |

Zimmer, Inc. is a leading orthopedic medical device manufacturer based in Warsaw, US. The company specializes in innovative surgical implants and trauma solutions. Zimmer, Inc. maintains a strong FDA 510(k) regulatory record with cleared devices from total submissions since 1976. Orthopedic devices represent approximately 90% of the company's submission portfolio. The company remains actively engaged in product development, with the latest FDA 510(k) clearance in 2026. Recent cleared devices reflect the company's focus on joint reconstruction and trauma fixation. Notable ...

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k081860/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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