

**K860356 ORTHOBLEND POWERED BONE MILL**Jul 8, 1986  
159 days to decisionK860356 · Product code: **LYS** · Orthopedic  
Source: <https://www.510kdatabase.net/k860356/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Bone Mill (LYS)                    |
| Date received         | Jan 30, 1986                       |
| Decision date         | Jul 8, 1986                        |
| Days to decision      | 159 days                           |
| Third-party review    | No                                 |

**APPLICANT**

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|----------------|---|
| Company        | <b>Zimmer, Inc.</b>   |
| Location       | Warsaw, IN, US  |
| Contact        | HENRY A QUELLO  |
| Website        | <a href="https://www.zimmerbiomet.com">https://www.zimmerbiomet.com</a> |
| 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 ...

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k860356/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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