

K760636 ORTHOMIXOct 5, 1976
22 days to decisionK760636 · Product code: **JDZ** · Orthopedic
Source: <https://www.510kdatabase.net/k760636/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixer, Cement, For Clinical Use (JDZ)
Date received	Sep 13, 1976
Decision date	Oct 5, 1976
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Diemolding Corp.
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1976-1996

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Device record: <https://www.510kdatabase.net/k760636/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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