FDA 510(k) Premarket Notification

K760636 ORTHOMIX

CLEARED

Oct 5, 1976
22 days to decision

K760636 · 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