FDA 510(k) Premarket Notification

K023360 LEFORTE SYSTEM BONE PLATE

CLEARED

Oct 25, 2002
18 days to decision

K023360 · Product code: **KTW** · Orthopedic
Source: https://www.510kdatabase.net/k023360/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Single Component (KTW)
Date received	Oct 7, 2002
Decision date	Oct 25, 2002
Days to decision	18 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Jeil Medical Corporation**
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	53 submissions · 53 cleared · 2002-2026