FDA 510(k) Premarket Notification

K052735 ALLOCRAFT DBM

CLEARED

Dec 28, 2005
89 days to decision

K052735 · Product code: **MBP** · Orthopedic
Source: https://www.510kdatabase.net/k052735/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Osteoinduction (w/o Human Growth Factor) (MBP)
Date received	Sep 30, 2005
Decision date	Dec 28, 2005
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lifecell Corp.**
Location	Washington, DC, US
Contact	HOWARD M HOLSTEIN
Website	http://www.lifecell.com/
510(k) history	10 submissions · 10 cleared · 2005-2013