

K052735 ALLOCRAFT DBMDec 28, 2005
89 days to decisionK052735 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k052735/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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