

**K073329 SYGNAL DBM**Feb 28, 2008  
93 days to decisionK073329 · Product code: **MBP** · Orthopedic  
Source: <https://www.510kdatabase.net/k073329/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Osteoinduction (w/o Human Growth Factor) (MBP)
Date received	Nov 27, 2007
Decision date	Feb 28, 2008
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Musculoskeletal Transplant Foundation</b>
Location	Edison, NJ, US
Contact	NANCY BENNEWITZ
510(k) history	24 submissions · 24 cleared · 2004-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073329/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 18, 2026