

**K130703 VIVORTE BVF**Sep 12, 2013  
181 days to decisionK130703 · Product code: **MBP** · Orthopedic  
Source: <https://www.510kdatabase.net/k130703/>**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	Mar 15, 2013
Decision date	Sep 12, 2013
Days to decision	181 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vivorte, Inc.</b>
Location	Fort Wayne, IN, US
Contact	STEPHEN J PEOPLES
510(k) history	4 submissions · 4 cleared · 2013-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130703/> 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 May 17, 2026