FDA 510(k) Premarket Notification

K130703 VIVORTE BVF

CLEARED

Sep 12, 2013
181 days to decision

K130703 · Product code: **MBP** · Orthopedic
Source: https://www.510kdatabase.net/k130703/

SUBMISSION DETAILS

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

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