

**K062833 POWERED EZ-IO BONE MARROW ASPIRATION SYSTEM**Nov 30, 2006  
70 days to decisionK062833 · Product code: **FCF** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k062833/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy, Mechanical, Gastrointestinal (FCF)
Date received	Sep 21, 2006
Decision date	Nov 30, 2006
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vidacare Corporation</b>
Location	Irvine, CA, US
Contact	GRACE HOLLAND
510(k) history	19 submissions · 19 cleared · 2004-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062833/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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