

**K081713 VERTEBRAL ACCESS SYSTEM BY VIDACARE**Nov 21, 2008  
157 days to decisionK081713 · Product code: **MOQ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k081713/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Battery, Replacement, Rechargeable (MOQ)
Date received	Jun 17, 2008
Decision date	Nov 21, 2008
Days to decision	157 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/k081713/> 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 7, 2026