

**K112468 THE ONCONTROL BONE ACCESS SYSTEM BY VIDACARE**

Oct 20, 2011  
55 days to decision

K112468 · Product code: **KNW** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k112468/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Aug 26, 2011
Decision date	Oct 20, 2011
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Regulatory Specialists, Inc.</b>
Location	Irvine, CA, US
Contact	GRACE HOLLAND
510(k) history	1 submissions · 1 cleared · 2011-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k112468/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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