

**K120189 BONETRAP**Feb 22, 2012  
30 days to decisionK120189 · Product code: **KNW** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k120189/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Jan 23, 2012
Decision date	Feb 22, 2012
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Berni Farns Incorporated</b>
Location	Fernandina Beach, FL, US
Contact	NICOLA BERNI
510(k) history	1 submissions · 1 cleared · 2012-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120189/> 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 15, 2026