

**K130282 REMINGTON MEDICAL CENTREFIRE 22 BIOPSY  
INSTRUMENT**May 13, 2013  
97 days to decisionK130282 · Product code: **KNW** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k130282/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Feb 5, 2013
Decision date	May 13, 2013
Days to decision	97 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Remington Medical, Inc.</b>
Location	Great Neck, NY, US
Contact	CAITLIN SENTER, MS, RAC
510(k) history	19 submissions · 19 cleared · 1993-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130282/> 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 May 25, 2026