

**K191315 Remington Medical, Inc. Automatic Cutting Needle (NAC)**

Jul 12, 2019  
58 days to decision

K191315 · Product code: **KNW** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k191315/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	May 15, 2019
Decision date	Jul 12, 2019
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Remington Medical, Inc.</b>
Location	Great Neck, NY, US
Contact	Caitlin Senter
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/k191315/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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