

**K173740 Reprocessed DePuy Mitek Ablation Wand**Mar 19, 2018  
102 days to decisionK173740 · Product code: **NUJ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k173740/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation Accessories, Laparoscopic & Endoscopic, Reprocessed (NUJ)
Date received	Dec 7, 2017
Decision date	Mar 19, 2018
Days to decision	102 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Renovo, Inc.</b>
Location	Bend, OR, US
Contact	Mark K. Wells
510(k) history	6 submissions · 6 cleared · 2017-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k173740/> 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 16, 2026