

**K133281 NEOTRACT UROLIFT SYSTEM**Dec 20, 2013  
56 days to decisionK133281 · Product code: **PEW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k133281/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Implantable Transprostatic Tissue Retractor System (PEW)
Date received	Oct 25, 2013
Decision date	Dec 20, 2013
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Neotract, Inc.</b>
Location	Pleasanton, CA, US
Contact	Nancy E Isaac
510(k) history	17 submissions · 16 cleared · 2006-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133281/> 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 26, 2026