

**K241700 Tenex 2nd Generation System**Nov 18, 2024  
158 days to decisionK241700 · Product code: **LFL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k241700/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Jun 13, 2024
Decision date	Nov 18, 2024
Days to decision	158 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Trice Medical, Inc.</b>
Location	King Of Prussia, PA, US
Contact	David Vancelette
510(k) history	3 submissions · 3 cleared · 2014-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k241700/> 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 13, 2026