

**K190281 Ultrasonic Surgical System**Oct 31, 2019  
262 days to decisionK190281 · Product code: **LFL** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k190281/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Feb 11, 2019
Decision date	Oct 31, 2019
Days to decision	262 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medisonic Technology CO</b>
Location	Las Vegas, NV, US
Contact	Reginald Bart-Williams
510(k) history	1 submissions · 1 cleared · 2019-2019

**REGULATORY CONSULTANT**

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Consulting firm	<b>FDA 510k Consultants, LLC</b>
Contact	John Gillespy

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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