

**K182211 Aurora Surgiscope System**Jan 18, 2019  
156 days to decisionK182211 · Product code: **GWG** · Neurology  
Source: <https://www.510kdatabase.net/k182211/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscope, Neurological (GWG)
Date received	Aug 15, 2018
Decision date	Jan 18, 2019
Days to decision	156 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Rebound Therapeutics Corporation</b>
Location	Irvine, CA, US
Contact	Jane Metcalf
510(k) history	2 submissions · 2 cleared · 2019-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k182211/> 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