

**K180653 Spartan eCoil System**Dec 4, 2018  
266 days to decisionK180653 · Product code: **HCG** · Neurology  
Source: <https://www.510kdatabase.net/k180653/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Neurovascular Embolization (HCG)
Date received	Mar 13, 2018
Decision date	Dec 4, 2018
Days to decision	266 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spartan Micro, Inc.</b>
Location	Freemont, CA, US
Contact	Mark Dias
510(k) history	3 submissions · 3 cleared · 2018-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180653/> 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 27, 2026