

**K053160 MICRUS MODIFIED MICROCOIL 18-SYSTEM,  
CERECYTE, MODEL CSP & CHE**Dec 7, 2005  
23 days to decisionK053160 · Product code: **HCG** · Neurology  
Source: <https://www.510kdatabase.net/k053160/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Device, Neurovascular Embolization (HCG)
Date received	Nov 14, 2005
Decision date	Dec 7, 2005
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Micrus Endovascular Corporation</b>
Location	Sunnyvale, CA, US
Contact	MARGARET WEBBER
510(k) history	23 submissions · 23 cleared · 2005-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k053160/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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