

**K032872 MICRUS MICROCOIL SYSTEM**Nov 28, 2003  
74 days to decisionK032872 · Product code: **HCG** · Neurology  
Source: <https://www.510kdatabase.net/k032872/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Neurovascular Embolization (HCG)
Date received	Sep 15, 2003
Decision date	Nov 28, 2003
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Micrus Corp.</b>
Location	Mountain View, CA, US
Contact	MARGARET WEBBER
510(k) history	6 submissions · 6 cleared · 2001-2004

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032872/> 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 June 30, 2026