

**K033372 SAPPHIRE DETACHABLE FIBERED COIL SYSTEM,
HELIX FIBERED, CYCLONE, MULTIPLE MODELS**Jan 9, 2004
79 days to decisionK033372 · Product code: **HCG** · Neurology
Source: <https://www.510kdatabase.net/k033372/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Neurovascular Embolization (HCG)
Date received	Oct 22, 2003
Decision date	Jan 9, 2004
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Micro Therapeutics, Inc.
Location	Aliso Viejo, CA, US
Contact	FLORIN TRUUVERT
510(k) history	51 submissions · 50 cleared · 1994-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k033372/> Data sourced from FDA 510(k) public records (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. - Generated June 14, 2026