

**K061063 MODIFICATION TO: SILVERHAWK PERIPHERAL
PLAQUE EXCISION SYSTEM, MODELS P4028/FG08300**May 18, 2006
31 days to decisionK061063 · Product code: **MCW** · Cardiovascular
Source: <https://www.510kdatabase.net/k061063/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Peripheral, Atherectomy (MCW)
Date received	Apr 17, 2006
Decision date	May 18, 2006
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Foxhollow Technologies, Inc.
Location	Redwood City, CA, US
Contact	MELISSA MURPHY
510(k) history	2 submissions · 2 cleared · 2006-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k061063/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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