

**K071826 THE RINSPIRATOR, MODELS FG09770, FG08776,
FG08777, FG08910**Nov 9, 2007
129 days to decisionK071826 · Product code: **QEZ** · Cardiovascular
Source: <https://www.510kdatabase.net/k071826/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Aspiration Thrombectomy Catheter (QEZ)
Date received	Jul 3, 2007
Decision date	Nov 9, 2007
Days to decision	129 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Fox Hollow Technologies
Location	Redwood City, CA, US
Contact	SRIDEVI SHESHADRI
510(k) history	5 submissions · 5 cleared · 2003-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k071826/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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