

K102439 FUNNEL GUIDE CATHETER MODEL G1020608Jan 12, 2012
504 days to decisionK102439 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k102439/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Aug 26, 2010
Decision date	Jan 12, 2012
Days to decision	504 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lazarus Effect, Inc.
Location	Felton, CA, US
Contact	CARRIE NEUBERGER
510(k) history	1 submissions · 1 cleared · 2012-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k102439/> 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 15, 2026