

**K072269 ANGIOJET ULTRA DVX THROMBECTOMY SET,
MODEL: 106552**Sep 8, 2007
24 days to decisionK072269 · Product code: **QEZ** · Cardiovascular
Source: <https://www.510kdatabase.net/k072269/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Aspiration Thrombectomy Catheter (QEZ)
Date received	Aug 15, 2007
Decision date	Sep 8, 2007
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Possis Medical, Inc.
Location	Walker, MI, US
Contact	MIKE BURNSIDE
510(k) history	34 submissions · 34 cleared · 1983-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072269/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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