FDA 510(k) Premarket Notification

K072269 ANGIOJET ULTRA DVX THROMBECTOMY SET, MODEL: 106552

CLEARED

Sep 8, 2007
24 days to decision

K072269 · 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