FDA 510(k) Premarket Notification

K111182 ANGIOJET SOLENT OMNI THROMBECTOMY SET

CLEARED

May 24, 2011
27 days to decision

K111182 · Product code: **QEZ** · Cardiovascular
Source: https://www.510kdatabase.net/k111182/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Aspiration Thrombectomy Catheter (QEZ)
Date received	Apr 27, 2011
Decision date	May 24, 2011
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medrad, Inc.**
Location	Mchenry, IL, US
Contact	Doug Atkins
Website	http://www.medrad.com/
510(k) history	105 submissions · 105 cleared · 1976-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k111182/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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