

**K982657 AMPLATZ THROMBECTOMY DEVICE MODELS ATD 601, ATD 602, ATD 801, ATD 802, FOOT PEDAL ASSEMBLY MODEL AK 100**

Nov 18, 1998  
111 days to decision

K982657 · Product code: **MCW** · Cardiovascular  
Source: <https://www.510kdatabase.net/k982657/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peripheral, Atherectomy (MCW)
Date received	Jul 30, 1998
Decision date	Nov 18, 1998
Days to decision	111 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Microvena Corp.</b>
Location	Findley, MN, US
Contact	TONYA WEIGEL
510(k) history	18 submissions · 18 cleared · 1990-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k982657/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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