

**K896096 SOLOSTICK (AMPLATZ DILATOR NEEDLE)**Apr 10, 1990  
172 days to decisionK896096 · Product code: **DRE** · CardiovascularSource: <https://www.510kdatabase.net/k896096/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Oct 20, 1989
Decision date	Apr 10, 1990
Days to decision	172 days
Third-party review	No

**APPLICANT**

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Company	<b>Microvena Corp.</b>
Location	Findley, MN, US
Contact	CONSTANCE G BUNDY
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/k896096/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 4, 2026