

**K901502 AMPLATZ GOOSE-NECK SNARE**Dec 3, 1990  
249 days to decisionK901502 · Product code: **DXE** · CardiovascularSource: <https://www.510kdatabase.net/k901502/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Mar 29, 1990
Decision date	Dec 3, 1990
Days to decision	249 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/k901502/>; 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