

**K895098 BIOPSY FORCEPS**Oct 30, 1989  
76 days to decisionK895098 · Product code: **DWZ** · CardiovascularSource: <https://www.510kdatabase.net/k895098/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biopsy, Endomyocardial (DWZ)
Date received	Aug 15, 1989
Decision date	Oct 30, 1989
Days to decision	76 days
Third-party review	No

**APPLICANT**

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Company	<b>Annex Medical, Inc.</b>
Location	Eden Prairie, MN, US
Contact	STUART LIND
510(k) history	23 submissions · 23 cleared · 1989-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895098/>; 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 1, 2026