

**K990809 CADISCOPE ELECTRONIC STETHOSCOPE**Jun 9, 1999  
90 days to decisionK990809 · Product code: **DQD** · CardiovascularSource: <https://www.510kdatabase.net/k990809/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Mar 11, 1999
Decision date	Jun 9, 1999
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Caditec AG</b>
Location	Mundelein, IL, US
Contact	BETTY LOCK
510(k) history	1 submissions · 1 cleared · 1999-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990809/> 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