

**K935732 CARDIO PERFECT ST 2001**Apr 4, 1996  
855 days to decisionK935732 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k935732/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Dec 1, 1993
Decision date	Apr 4, 1996
Days to decision	855 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cardio Control BV</b>
Location	The Netherlands, NL
Contact	DICK F VAN LUIJK
510(k) history	4 submissions · 4 cleared · 1992-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k935732/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated July 4, 2026