

K925439 MICROSNARESep 24, 1993
331 days to decisionK925439 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k925439/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Oct 28, 1992
Decision date	Sep 24, 1993
Days to decision	331 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Microvena Corp.
Location	Findley, MN, US
Contact	MICHAEL RENNER
510(k) history	18 submissions · 18 cleared · 1990-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925439/> Data sourced from FDA 510(k) public records (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. - Generated July 4, 2026