FDA 510(k) Premarket Notification

K925439 MICROSNARE

CLEARED

Sep 24, 1993
331 days to decision

K925439 · 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