FDA 510(k) Premarket Notification

K991898 ULTRA-SELECT GUIDEWIRE, HYTEK GUIDEWIRE

CLEARED

Jun 16, 1999
12 days to decision

K991898 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k991898/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 4, 1999
Decision date	Jun 16, 1999
Days to decision	12 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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