FDA 510(k) Premarket Notification

K042685 METRICATH SYSTEM

CLEARED

Nov 18, 2004
50 days to decision

K042685 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k042685/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Sep 29, 2004
Decision date	Nov 18, 2004
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Angiometrx, Inc.**
Location	Seattle, WA, US
Contact	TIM VERSPAGEN
510(k) history	3 submissions · 3 cleared · 2003-2005