FDA 510(k) Premarket Notification

K083922 MICRUS MICROCATHETER, COURIER FLEX, MODEL MSF 170000-00

CLEARED

May 6, 2009
126 days to decision

K083922 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k083922/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Dec 31, 2008
Decision date	May 6, 2009
Days to decision	126 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Micrus Endovascular Corporation**
Location	Sunnyvale, CA, US
Contact	PATRICK LEE
510(k) history	23 submissions · 23 cleared · 2005-2011