FDA 510(k) Premarket Notification

CLEARED

K102818 MICRUS ONE 2 ONE GUIDEWIRE, MICRUS ONE 2 ONE SOFT GUIDEWIRE

Dec 3, 2010
66 days to decision

K102818 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k102818/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Sep 28, 2010
Decision date	Dec 3, 2010
Days to decision	66 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