

**K102818 MICRUS ONE 2 ONE GUIDEWIRE, MICRUS ONE 2 ONE
SOFT GUIDEWIRE**Dec 3, 2010
66 days to decisionK102818 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k102818/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 19, 2026