

K090274 MICRUS ONE 2 ONE GUIDEWIRE AND SOFT GUIDEWIREMar 3, 2009
27 days to decisionK090274 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k090274/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 4, 2009
Decision date	Mar 3, 2009
Days to decision	27 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/k090274/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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