

K060010 NAUTIFLUXAug 4, 2006
213 days to decisionK060010 · Product code: **DQO** · Cardiovascular
Source: <https://www.510kdatabase.net/k060010/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jan 3, 2006
Decision date	Aug 4, 2006
Days to decision	213 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sedat
Location	San Diego, CA, US
Contact	LAETITIA COUSIN
510(k) history	3 submissions · 3 cleared · 2006-2008

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