

K021323 MODIFICATION TO SAFE-STEER GUIDE WIRE SYSTEMMay 22, 2002
26 days to decisionK021323 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k021323/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Apr 26, 2002
Decision date	May 22, 2002
Days to decision	26 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Intra Luminal Therapeutics, Inc.
Location	Carlsbad, CA, US
Contact	PAMELA MISAJON
510(k) history	20 submissions · 20 cleared · 2000-2005

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