

**K011986 SAFE-STEER GUIDE WIRE SYSTEM**Apr 3, 2002  
281 days to decisionK011986 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k011986/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 26, 2001
Decision date	Apr 3, 2002
Days to decision	281 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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