

**K060302 INSCOPE PRECISION BALLOON DILATOR - WIRE  
GUIDED**Apr 11, 2006  
64 days to decisionK060302 · Product code: **KNQ** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k060302/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Esophageal (KNQ)
Date received	Feb 6, 2006
Decision date	Apr 11, 2006
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cordis Europa, N.V.</b>
Location	Warren, NJ, US
Contact	HARM HOVINGA
510(k) history	7 submissions · 5 cleared · 2001-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k060302/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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