

**K051860 FLEXIBLE INTRODUCER CANNULA, MODEL 135-1837**Aug 22, 2005  
45 days to decisionK051860 · Product code: **MIA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k051860/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Spinal, Short Term (MIA)
Date received	Jul 8, 2005
Decision date	Aug 22, 2005
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Epimed International, Inc.</b>
Location	Johnstown, NY, US
Contact	CHRISTOPHER LAKE
510(k) history	14 submissions · 14 cleared · 1998-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051860/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 23, 2026