

**K060235 ENDO-EASE ENDOSCOPIC OVERTUBE**Aug 4, 2006  
186 days to decisionK060235 · Product code: **FDA** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k060235/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enteroscope And Accessories (FDA)
Date received	Jan 30, 2006
Decision date	Aug 4, 2006
Days to decision	186 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Spirus Medical, Inc.</b>
Location	Ayer, MA, US
Contact	PAMELA PAPINEAU
510(k) history	5 submissions · 5 cleared · 2005-2008

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k060235/> 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 25, 2026