

K091783 THIRD EYE RETROSCOPE AUXILIARY ENDOSCOPE SYSTEMJul 7, 2009
20 days to decisionK091783 · Product code: **FDF** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k091783/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Colonoscope And Accessories, Flexible/rigid (FDF)
Date received	Jun 17, 2009
Decision date	Jul 7, 2009
Days to decision	20 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Avantis Medical Systems, Inc.
Location	Sunnyvale, CA, US
Contact	AMRITA SETHI
510(k) history	6 submissions · 6 cleared · 2007-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091783/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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