

**K093037 PENTEX EC-3890LI**Dec 28, 2009  
90 days to decisionK093037 · Product code: **FDF** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k093037/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Colonoscope And Accessories, Flexible/rigid (FDF)
Date received	Sep 29, 2009
Decision date	Dec 28, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pentax Medical Company</b>
Location	Montvale, NJ, US
Contact	LISA LANCIA
510(k) history	12 submissions · 12 cleared · 2004-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093037/> 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 May 17, 2026