

**K161222 PENTAX MEDICAL ED-3490TK, Video Duodenoscope**Feb 7, 2018  
649 days to decisionK161222 · Product code: **FDT** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k161222/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Duodenoscope And Accessories, Flexible/rigid (FDT)
Date received	Apr 29, 2016
Decision date	Feb 7, 2018
Days to decision	649 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pentax Medical</b>
Location	Montvale, NJ, US
Contact	KANCHANA IYER
510(k) history	6 submissions · 6 cleared · 2013-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k161222/> 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 28, 2026