

**K093106 ENDOSCOPE REPROCESSOR OER-PRO**Feb 12, 2010  
134 days to decisionK093106 · Product code: **FEB** · General Hospital  
Source: <https://www.510kdatabase.net/k093106/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Cleaning, For Endoscope (FEB)
Date received	Oct 1, 2009
Decision date	Feb 12, 2010
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Olympus Medical Systems Corporation</b>
Location	Melville, NY, US
Contact	LAURA STORMS-TYLER
510(k) history	81 submissions · 81 cleared · 2004-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093106/> 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 14, 2026