

**K091369 ELECTROFLUIDGRAPH**Oct 14, 2009  
159 days to decisionK091369 · Product code: **MNW** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k091369/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Body Composition (MNW)
Date received	May 8, 2009
Decision date	Oct 14, 2009
Days to decision	159 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Akern USA, LLC</b>
Location	Colorado, CO, US
Contact	KEVIN WELLS
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091369/> 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 22, 2026