

**K092090 ERBELIFT HAND PUMP AND FLEXIBLE PROBE**Nov 30, 2009  
139 days to decisionK092090 · Product code: **FBK** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k092090/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Injection Needle, Gastroenterology-urology (FBK)
Date received	Jul 14, 2009
Decision date	Nov 30, 2009
Days to decision	139 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Byrne Medical, Inc.</b>
Location	Houston, TX, US
Contact	CHRIS HIERHOLZER
510(k) history	12 submissions · 12 cleared · 2003-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092090/> 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 14, 2026