

**K904849 ORIGINEEDLE(TM)**Mar 29, 1991  
154 days to decisionK904849 · Product code: **FHO** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k904849/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pneumoperitoneum Needle (FHO)
Date received	Oct 26, 1990
Decision date	Mar 29, 1991
Days to decision	154 days
Third-party review	No

**APPLICANT**

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Company	<b>Origin Medsystems, Inc.</b>
Location	San Mateo, CA, US
Contact	RICHARD RUSH
510(k) history	50 submissions · 49 cleared · 1989-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904849/>; 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 22, 2026